



- 17 -
Patent
SYR-DPP-IV-5004-C3

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re the Application of:) Group Art Unit: 1614
Jun Feng et al.) Examiner: Not Yet Assigned
Serial No.: 10/809,638)
Filed: March 24, 2004)
For: DIPEPTIDYL PEPTIDASE)
INHIBITORS)

INFORMATION DISCLOSURE STATEMENT

Mail Stop Amendment
Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

Sir:

In accordance with 37 CFR §§ 1.97 and 1.98, the items identified in this Information Disclosure Statement ("IDS") are brought to the attention of the Office. The items are listed on the attached form PTO-1449, and copies are included for the Examiner's convenience. As the Office no longer requires copies of U.S. patents and applications, the U.S. copies are not being submitted with this IDS. However, if the Examiner would like copies of the U.S. cited references, Applicants will provide the references upon request. The item listed on the attached form PTO-1449 at "EN" is a non-English language article. In accordance with 37 CFR § 1.98(a)(3)(i), the following is a concise explanation of the relevance of this article:

The article by P.O. Bezuglyi relates to the synthesis of arylsulfonyl hydrazides of 3-R-quinazolone-4-carbonyl-2-acid.

The items identified in this IDS may or may not be "material" pursuant to 37 CFR § 1.56. The submission thereof by Applicant is not to be construed as an admission that any such patent, publication or other information referred to therein is material or considered to be material (37 CFR § 1.97(h)), or even qualifies as "prior art" under 35 USC § 102 with respect to this invention unless specifically designated by Applicants as such.

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Substitute for form 1449A/PTO				Complete if Known	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT <i>(use as many sheets as necessary)</i>				Application Number	10/809,638
				Filing Date	March 24, 2004
				First Named Inventor	Jun Feng
				Group Art Unit	1614
				Examiner Name	Not Yet Assigned
Sheet	1	of	10	Attorney Docket Number	SYR-DPP-IV-5004-C3

U.S. PATENT DOCUMENTS					
Examiner Initials *	Cite No. ¹	Document Number	Publication Date/ Issue Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number - Kind Code ² (if known)			
AA		US1974/3823135	07-09-1974	Pilgram et al.	
AB		US1996/5512549	04-30-1996	Chen et al.	
AC		US1996/5580979	12-03-1996	Bachovchin	
AD		US1997/5614492	03-25-1997	Habener	
AE		US2000/6156739	12-5-2000	Griffin et al.	
AF		US2000/6166063	12-26-2000	Villhauer	
AG		US2001/6258597-B1	07-10-2001	Bachovchin	
AH		US2001/0020006-A1	09-06-2001	Demuth et al.	
AI		US2001/6303661-B1	10-16-2001	Demuth et al.	
AJ		US2001/6319893-B1	11-20-2001	Demuth et al.	
AK		US2001/0051646-A1	12-13-2001	Demuth et al.	
AL		US2002/0049153-A1	04-25-2002	Bridon et al.	
AM		US2002/0049164-A1	04-25-2002	Demuth et al.	
AN		US2002/6380398-B2	04-30-2002	Kanstrup et al.	
AO		US2002/0082427-A1	06-27-2002	Demuth et al.	
AP		US2002/6448045-B1	09-10-2002	Levine et al.	
AQ		US2002/0198242-A1	12-26-2002	Demuth et al.	
AR		US2002/0198380-A1	12-26-2002	Belzer et al.	
AS		US2002/6500804-B2	12-31-2002	Demuth et al.	
AT		US2003/0008925-A1	01-09-2003	Demuth et al.	
AU		US2003/6548481-B1	04-15-2003	Demuth et al.	
AV		US2003/0092630-A2	05-15-2003	Demuth et al.	
AW		US2003/0119750-A1	06-26-2003	Demuth et al.	
AX		US2003/0130199-A1	07-10-2003	von Hoersten et al.	
AY		US2003/0134802-A1	07-17-2003	Demuth et al.	
AZ		US2003/0135023-A1	07-17-2003	Demuth et al.	
BA		US2003/0148961-A1	08-07-2003	Heiser et al.	
BB		US2003/0153509-A1	08-14-2003	Bachovchin et al.	
BC		US2003/0162820-A1	08-28-2003	Demuth et al.	
BD		US2003/0166578-A1	09-04-2003	Arch et al.	
BE		US2003/6620910-B1	09-16-2003	Calas et al.	
BF		US2003/0176357-A1	09-18-2003	Pospisilik et al.	
BG		US2003/0199451-A1	10-23-2003	Mogensen et al.	
BH		US2003/0199672-A1	10-23-2003	Knudsen et al.	

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1 Applicant's unique citation designation number (optional). 2 Applicant is to place a check mark here if English language Translation is attached.
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				First Named Inventor	Jun Feng
				Group Art Unit	1614
				Examiner Name	Not Yet Assigned
Sheet	2	of	10	Attorney Docket Number	SYR-DPP-IV-5004-C3

BI	US2003/0236272-A1	12-25-2003	Richard David Carr	
BJ	US2004/6703238-B2	03-09-2004	Bachovchin	
BK	US2004/0054171-A1	03-18-2004	Jensen et al.	
BL	US2004/0058876-A1	03-25-2004	Hoffmann et al.	
BM	US2004/0132732-A1	07-08-2004	Han et al.	
BN	US2004/0167191-A1	08-26-2004	Demuth et al.	
BO	US2004/0171555-A1	09-02-2004	Demuth et al.	

FOREIGN PATENT DOCUMENTS						
Examiner Initials*	Cite No. ¹	Foreign Patent Document	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T ⁶
		Country Code ³ - Number ⁴ - Kind Code ⁵ (if known)				
BP		FR 2.162.106 (English Abstract-1973)	11-30-1972	Amschler et al.		
BQ		WO 89/10701	11-16-1989	BASF		
BR		EP 0378255-A2	07-18-1990	Janssen Pharmaceutica		
BS		GB 2230527-A	10-24-1990	Imperial Chemical Industries Plc		
BT		WO 91/12001	08-22-1991	Merck & Co., Inc.		
BU		WO 93/21162	01-28-1993	Nissan Chemical Industries, Ltd.		
BV		WO 93/08259 (A2)	04-29-1993	New England Medical Center Hospitals, Inc.		
BW		WO 93/08259 (A3)	04-29-1993	New England Medical Center Hospitals, Inc.		
BX		EP 0547442-A1	06-23-1993	E.R. Squibb & Sons, Inc.		
BY		WO 94/03055	02-17-1994	U.S. Government, Secty. HHS		
BZ		EP 0587377-A2	03-16-1994	Eli Lilly and Company		
CA		WO 95/35031	12-28-1995	La Trobe University		
CB		WO 96/32384	10-17-1996	Taiho Pharmaceutical Co., Ltd.		
CC		WO 96/38550	12-05-1996	Dana-Farber Cancer Institute, Inc.		
CD		WO 97/40832	11-06-1997	Hans-Knoll-Institut Fur Naturstoff		
CE		JP 9295977	11-18-1997	Terumo Corp.		
CF		WO 98/00439	01-08-1998	Trustees of Tufts College		
CG		WO 98/24780	06-11-1998	Amgen Inc.		
CH		WO 99/16864	04-08-1999	Point Therapeutics, Inc.		
CI		WO 99/38501	08-05-1999	Trustees of Tufts University		

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Sheet	3	of	10	Attorney Docket Number	SYR-DPP-IV-5004-C3

	CJ	WO 99/50249	10-07-1999	Du Pont Pharmaceuticals Company	
	CK	WO 99-61431	12-02-1999	Probiodrug	
	CL	WO 99/67278	12-29-1999	Pro-Biodrug	
	CM	WO 99/67279	12-29-1999	Pro-Biodrug	
	CN	WO 00/07617	02-17-2000	Novo Nordisk	
	CO	WO 00/09666	02-24-2000	U.S. Government, Secty. HHS	
	CP	WO 00/15211	03-23-2000	Akesis Pharmaceuticals, Inc.	
	CQ	WO 00/76986-A1	04-11-2000	Probiodrug	
	CR	WO 00/34241	06-15-2000	Novartis AG	
	CS	WO 00/47219	08-17-2000	Ontogeny, Inc.	
	CT	WO 00/53171	09-14-00	Molteni L. E C. Dei Fratelli Alitti Societa' Di Esercizio S.P.A.	
	CU	WO 00/57721	10-05-2000	Akesis Pharmaceuticals, Inc.	
	CV	WO 01/14318-A2	03-01-2001	Probiodrug	
	CW	WO 01/34594-A1	05-17-2001	Guilford Pharmaceuticals, Inc.	
	CX	WO 01/52825-A2	07-26-2001	Novartis AG	
	CY	WO 01/56988-A1	08-09-2001	Kirin Beer Kabaushiki Kaisha	
	CZ	WO 01/70729-A1	09-27-2001	Sanofi-Synthelabo	
	DA	WO 01/97808-A1	12-27-2001	Smithkline Beecham PLC	
	DB	WO 02/34242-A2	05-02-2002	Probiodrug AG	
	DC	WO 02/34243-A2	05-02-2002	Probiodrug AG	
	DD	WO 02/083109-A1	10-24-2002	Ferring BV	
	DE	JP 2002/338466	11-27-2002	Tanabe Seiyaku Co Ltd	
	DF	WO 03/002593-A2	01-09-2003	Probiodrug AG	
	DG	WO 03/002595-A2	01-09-2003	Probiodrug AG	
	DH	WO 03/002596-A2	01-09-2003	Probiodrug AG	
	DI	WO 03/016335-A2	02-27-2003	Probiodrug AG	
	DJ	WO 03/022871-A2	03-20-2003	Probiodrug AG	
	DK	WO 03/026652-A1	04-03-2003	Bristol-Myers Squibb Company	
	DL	WO 03/030946-A1	04-17-2003	Novartis AG	
	DM	WO 03/033524-A2	04-24-2003	Probiodrug AG	
	DN	JP 2003/128551	05-08-2003	Sankyo Co LTD	
	DO	WO 03/040174-A2	05-15-2003	Probiodrug AG	
	DP	WO 03/045228-A2	06-05-2003	Trustees of Tufts College	
	DQ	WO 03/045977-A2	06-05-2003	Trustees of Tufts College	
	DR	WO 03/048081-A2	06-12-2003	Bristol-Myers Squibb Company	
	DS	WO 03/048158-A1	06-12-2003	Bristol-Myers Squibb Company	
	DT	WO 03/057200-A2	07-17-2003	Novo Nordisk	

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Sheet	4	of	10	Attorney Docket Number	SYR-DPP-IV-5004-C3

	DU	WO 03/063903-A2	08-07-2003	Probiodrug AG	
	DV	WO 03/072556-A1	09-04-2003	Probiodrug AG	
	DW	WO 03/082898-A2	10-09-2003	Probiodrug AG	
	DX	WO 03/092605-A2	11-13-2003	Trustees of Tufts College	
	DY	WO 03/099279-A1	12-04-2003	Novartis AG	
	DZ	WO 03/099818-A1	12-04-2003	Chiron Corporation	
	EA	WO 03/106416-A2	12-24-2003	Smithkline Beecham Corporation	
	EB	WO 2004/017989-A1	03-04-2004	Probiodrug AG	
	EC	JP 2004/99600-A	04-02-2004	Sankyo Co. Ltd.	
	ED	WO 2004/031374-A2	04-15-2004	Probiodrug AG	
	EE	JP 2004/123738-A	04-22-2004	Takeda Chem Ind Ltd	
	EF	WO 2004/037176-A2	05-06-2004	Bristol-Myers Squibb Company	

OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS				
Examiner Initials *	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.		T ²
	EG	ARGAUD, DORIANE et al., Metformin decreases gluconeogenesis by enhancing the pyruvate kinase flux in isolated rat hepatocytes, European J. Biochem. 213, 1341-1348 (1993).		
	EH	ASHCROFT, STEPHEN J.H. et al., Structure-activity relationships of alloxan-like compounds derived from uric acid, Br. J. Pharmac. (1986), 89 pp. 469-472.		
	EI	BAL, GUNTHER, Dipeptidyl Peptidase IV and Prolyl Oligopeptidase: Design, Synthesis and Evaluation of Substrates and Inhibitors, (2002) Universiteit Antwerpen.		
	EJ	BARAKAT, S.E.S., Synthesis and hypoglycemic activity of some new 3-[4- [(cyclohexylamino) carbonyl] amino]sulfonylphenyl]-4(3H)-quinazolinones, Az. J. Pharm. Sci., Vol. 25, (2000), pp. 48-57.		
	EK	BARAKAT, S.E.S., Synthesis and Hypoglycemic Activity of Some New 4(3H) -Quinazolinone Analogues, Saudi Pharmaceutical Journal, Vol. 8, No.4 (2000) pp. 198-204.		
	EL	BAKER, B.R. et al., Irreversible Enzyme Inhibitors. On the Mode of Pyrimidine Binding of 5-alkyl and 5-Arylpyrimidines to Dihydrofolic Reductase (1,2), Journal of Heterocyclic Chemistry Vol. 4 (1967) pp. 39-48.		
	EM	BELGODERE, ELENA et al., Synthesis of Substituted Pyrimidines, Study of the Structure and of the Tautomeric Equilibria, (1976) Chem. Abstracts, Columbus, OH Vol. 85 No. 9.		
	EN	BEZUGLYI, P.O. et al., Synthesis of arylsulfonyl hydrazide of 3-R-quinazolone-4-carbonyl-2-acid, Pharmaceutical Journal (1979), pp. 70-71.		

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Sheet	5	of	10	Attorney Docket Number	SYR-DPP-IV-5004-C3

EO	BHADURI, A.P. et al., Urinary Metabolite of 2-Piperazine-3 (H)-4-Quinazolone (Centipiperalone), A Potent Blood Sugar Lowering Agent, Indian J. Biochem. Biophys., Vol. 12 (1975), pp. 413-414.
EP	BOURAS, MOHAMMED, et al., Metabolism of enterostatin in rat intestine, brain, membranes and serum: differential involvement of proline-specific peptidases, Peptides, Vol. 16, No. 3, (1995), pp. 399-405.
EQ	BRUN, JEAN-FREDERIC, et al., Effects of Oral Zinc Gluconate on Glucose Effectiveness and Insulin Sensitivity in Humans, Biological Trace Element Research Vol. 47 (1995), pp. 385-391.
ER	BUCKLEY, DI, Analysis of the Degradation of Insulinotropin [GLP-1 (7-37)] In Human Plasma and Production of Degradation Resistant Analogs.
ES	CHATTERJEE, A.K. et al., Effect of Centipiperalone in Insulin Deficient Diabetes, Indian Journal of Experimental Biology Vol. 18 (1980), pp. 1005-1008.
ET	CHATTERJEE, A.K. et al., Effect of Centipiperalone, a New Hypoglycemic Agent on Insulin Biosynthesis & Release from Isolated Pancreatic Islets of Rat, Indian Journal of Experimental Biology Vol. 20 (1981) pp.270-272.
EU	COPPOLA, GARY M. et al., 1-Aminomethylisoquinoline-4-carboxylates as Novel Dipeptidylpeptidase IV Inhibitors, Bioorganic & Medicinal Chemistry Letters Vol. 10 (2000), pp. 1555-1558.
EV	DEACON, CAROLYN F. et al., Degradation of Glucagon-Like Peptide 1 <i>in Vitro</i> Yields an N-Terminally Truncated Peptide That is a Major Endogenous Metabolite <i>in Vivo</i> , Journal of Clinical Endocrinology and Metabolism Vol. 80, No. 3 (1995), pp. 952-957.
EW	DEACON, CAROLYN F. et al., Both Subcutaneously and Intravenously Administered Glucagon-Like Peptide I Are Rapidly Degraded From the NH ₂ -Terminus in Type II Diabetic Patients and in Healthy Subjects, Diabetes, Vol. 44 (1996), pp. 1125-1131.
EX	DEACON, CAROLYN F. et al., Dipeptidyl peptidase IV Inhibition Influences GLP-1 Metabolism <i>in Vivo</i> , Regulatory Peptides Vol. 64 Issues 1-3 (1996) p.30.
EY	DEACON, CAROLYN F. et al., Dipeptidyl peptidase IV Inhibition Potentiates the Insulinotropic Effect of Glucagon-Like Peptide 1 in the Anesthetized Pig, Diabetes, Vol. 47 (1998), pp. 764-769.
EZ	DEACON, CAROLYN F. et al., Dipeptidyl peptidase IV Inhibition as an Approach to the Treatment and Prevention of Type 2 Diabetes: a Historical Perspective, Biochemical and Biophysical Research Communications 294 (2002), pp. 1-4.
FA	DEMUTH, HANS-ULRICH et al., Rebuttal to Deacon and Holst: "Metformin effects on depeptidyl peptidase IV degradation of glucagons-like peptide-1" versus "dipeptidyl peptidase inhibition as an approach to the treatment and prevention of type 2 diabetes: a historical perspective" Biochemical and Biophysical Research Communications 296 (2002) pp. 229-232.
FB	ENGEL, MICHAEL et al., The crystal structure of dipeptidyl peptidase IV (CD26) reveals its functional regulation and enzymatic mechanism, Proc. Nat. Acad. Sci. Early Edition (2003), pp. 1-6.
FC	FANTUS, I. GEORGE, et al., Mechanism of Action of Metformin: Insulin Receptor and Postreceptor Effects <i>in Vitro</i> and <i>in Vivo</i> , J. Clinical Endocrinology & Metabolism (1986), pp. 898-905.

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	FD	GARRATT, PETER J. et al., A Novel Synthesis of Dihydropyrimidines, J. Chem. Soc., Chem. Commun. (1987), pp.568-569.	
	FE	GARRATT, PETER J. et al., One-Carbon Compounds as Synthetic Intermediates. The Synthesis of Hydopyrimidines and Hydroquinazolines by Sequential Nucleophilic Addition to Diphenyl Cyanocarbonimidate With Concomitant Cyclization, J. Org. Chem. (1988), pp. 1062-1069.	
	FF	GAZIT, AVIV et al., Typhostins IV – Highly Potent Inhibitors of EGF Receptor Kinase. Structure-Activity Relationship Study of 4- Anilidoquinazolines, Bioorganic & Medicinal Chemistry, Vol. 4, No.8 (1996) pp. 1203-1207.	
	FG	GUERRIERI, N., et al., Vanadium Inhibition of Serine and Cysteine Proteases, Comparative Biochemistry and Physiology Part A 122 (1997), pp.331-336.	
	FH	GUPTA, C.M. et al., Drugs Acting on the Central Nervous System. Syntheses of Substituted Quinazolones and Quinazolines and Triazepino-and Triazoclonquinazolones, Division of Medicinal Chemistry, Central Drug Research Institute, Lucknow, India (1968), pp. 392-395.	
	FI	GUPTA, C.M. et al., New Potent Blood Sugar Lowering Compound, Nature, Vol. 223 (1969), p. 524.	
	FJ	GUPTA, C.M. et al., A Novel Class of Hypoglycaemic Agents: Syntheses & SAR in 2-Substituted 4(3H)-Quinazolones, 2-Substituted 4-Hydroxypolyethylene 5,6-pyrimidines & 3-Substituted 4-Oxo-pyrido [1,2-a] pyrimidines, Indian Journal of Chemistry, Vol. 9 (1971), pp. 201-206.	
	FK	HERMECZ, ISTVAN et al., Pyrido[1,2-a]Pyrimidines; New Chemical Entities in Medicinal Chemistry, Medicinal Research Reviews, Vol. 8, No. 2 (1988) pp.203-230.	
	FL	HINKE, SIMON A. et al., Metaformin Effects on Dipeptidylpeptidase IV Degradation of Glucagon-like Peptide-1, Biochemical and Biophysical Research Communications, 291 (2002) pp. 1302-1308.	
	FM	HINKE, SIMON A. et al., On Combination Therapy of Diabetes With Metaformin and Dipeptidyl Peptidase IV Inhibitors, Diabetes Care, Vol. 25, No. 8 (2002) pp. 1490-1492.	
	FN	HOLZ, GEORGE G. et al, Pancreatic Beta-Cells are Rendered Glucose-Competent by the Insulinotropic Hormone Glucagon-Like Peptide-1(7-37), Nature, Vol. 361 (1993), pp. 362-365.	
	FO	KHALID, NORAINI M., et al., Purification and Partial Characterization of a Prolyl-Dipeptidyl Aminopeptidase From Lactobacillus helveticus CNRZ 32, Applied and Environmental Microbiology (1990), pp. 381-388.	
	FP	KIEFFER, TIMOTHY J. et al., Degradation of Glucose-Dependant Insulinotropic Polypeptide and Truncated Glucagon-Like Peptide 1 in Vitro and in Vivo by Dipeptidyl Peptidase IV, Endocrinology, Vol. 136, No. 8 (1995) 3585-3596.	
	FQ	KIMURA, TOSHIKIRO et al., Oral Administration of Insulin as Poly(Vinyl Alcohol)-Gel Spheres in Diabetic Rats, Biological & Pharmaceutical Bulletin, Vol. 19, No. 6 (1996), 897-900.	
	FR	KOREEDA, YUJI et al., Isolation and Characterization of Dipeptidyl Peptidase IV From <i>Prevotella loescheii</i> ATCC 15930, Archives of Oral Biology, 46 (2001), 759-766.	

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Sheet	7	of	10	Attorney Docket Number	SYR-DPP-IV-5004-C3

FS	KUSAR, MIHAEL et al., Diethyl N,N-Dimethylaminomethylenemalonate in the Synthesis of Fused Heterocyclic Systems, <i>Heterocyclic Chem.</i> 33 (1996) pp. 1041-1046.
FT	LI JINPING, et al., Permolybdate and Pertungstate—Potent Stimulators of Insulin Effects in Rat Adipocytes: Mechanism of Action, <i>Biochemistry</i> , 34 (1995) 6218-6225.
FU	LIN, JIAN, Total Synthesis and Biological Evaluation of Fluoroolefin-containing Dipeptidyl Isosteres as Inhibitors of Dipeptidyl Peptidase IV (CD26), Dissertation presented to State University of New York at Albany, Department of Chemistry (1998).
FV	LOESER, ERIC et al., Selective N-Alkylation of Primary Amines with Chloroacetamides Under pH-Controlled Aqueous Conditions, <i>Synthetic Communications</i> , 32(3) (2002) pp. 403-409.
FW	MANNUCCI, EDUARDO, et al., Effect of Metformin on Glucagon-Like Peptide-1 (GLP-1) and Leptin Levels in Obese Nondiabetic Subjects, <i>Diabetes Care</i> , Vol. 24, No. 3 (2001) 489-494.
FX	MENTLEIN, ROLF et al., Dipeptidyl-Peptidase IV Hydrolyses gastric Inhibitory Polypeptide, Glucagon-Like Peptide-1(7-36)amide, Peptide Histidine Methionine and is Responsible for Their Degradation in Human Serum, <i>Eur. J. Biochem.</i> , Vol. 214, 829-835 (1991).
FY	MEYEROVITCH, JOSEPH et al., Oral Administration of Vanadate Normalizes Blood Glucose Levels in Streptozotocin-Treated Rats, <i>The Journal of Biological Chemistry</i> , Vol. 262, No. 14 (1987) 6658-6662.
FZ	MALLOY, J. ARDILL et al., Effect of Metformin Treatment on Gastric Acid Secretion Gastrointestinal Hormone Levels in Normal Subjects, <i>Diabetologia</i> , Vol. 19 (1980) 93-96.
GA	MUKERJEE, S.S. et al., Effect of 2-piperazino-4(3H)-quinazolinone monoacetate on the tissue respiration, glucose uptake and lactic acid production by rat hemidiaphragm, <i>Biochemical Pharmacology</i> , Vol. 23 (1974) 3066-3067.
GB	MUKERJEE, S.S. et al., Studies on the Mechanism of Centipiperalone-Induced Hypoglycemia, <i>Acta Diabet. Lat</i> 13, 8 (1976) p 8.
GC	MUKERJEE, S.S. et al., Chronic Toxicity Studies of a Hypoglycemic Compound: Centipiperalone in Rats & Rhesus Monkeys, <i>Indian Journal of Experimental Biology</i> , Vol. 17 (1979) pp. 1346-1349.
GD	MUKERJEE, S.S. et al., Tissue Distribution of [³ H]Centipiperalone after Oral Administration, <i>Indian J. Biochem. Biophys.</i> , Vol. 17 (1980) pp. 399-401.
GE	MUKHERJEE, SURATH K. et al., A novel hypoglycemic compound, <i>Biochemical Pharmacology</i> , Vol. 22 (1972) pp. 1529-1531.
GF	MUKHERJEE, SURATH K. et al., Effect of 2-piperazino-4(3H)-quinazolinone monoacetate on some aspects of carbohydrate metabolism of albino rats, <i>Biochemical Pharmacology</i> , Vol. 22 (1973) pp. 2205-2206.
GG	MUKHERJEE, SURATH K. et al., Studies on the Metabolic Changes Induced by a Synthetic Insulinogenic Agent, <i>Ind. J. Physiol. & Allied Sci.</i> , Vol. 30, No. 3 (1976) pp. 105-116.

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	GH	MUKHERJEE, SURATH K. et al., Influence of Timing Oral Dosing of a Novel Hypoglycaemic Agent A-4166 in Relation to Food, Diabetologia Vol. 38 A194 Supplement 1 (1995).	
	GI	MUKHERJEE, Subal S. et al., Studies on the Mechanism of Centipiperalone-Induced Hypoglycemia, Acta Diabet. Lat. 13, 8, (1976) pp. 8-19.	
	GJ	MURTHY, G. RAMA et al., New Hypoglycemic Agents: Part V – Synthesis & Hypoglycemic Activity of Some New 1-[[(p-(4-OXO-2-Methyl/Phenyl-3 (4H)-Quinazolinyl) Phenyl] 3-Aryl-2-Ureas, Indian Drugs, 25 (1) (1987) pp. 19-22.	
	GK	MURTHY, G. RAMA et al., New Hypoglycemic Agents: Synthesis and Hypoglycemic Activity of Some New 1-[{p-(4-OXO-2-Substituted-3(4H)-Quinazolinyl)-Phenyl]-3-Aryl/Cyclohexyl-2-Thioureas, Current Science, Vol. 56, No. 24 (1987) pp. 1263-1265.	
	GL	NAKAMURA, SEIJI, et al., Effect of Chronic Vanadate Administration in Partially Depancreatized Rats, Diabetes Research and Clinical Practice 27 (1995) pp. 51-59. (Abstract Only)	
	GM	OHKUBO, I., et al., Dipeptidyl Peptidase IV From Porcine Seminal Plasma: Purification, Characterization, and N-Terminal Amino Acid Sequence, J. Biochem. (Tokyo) (1994) 116(5) pp. 1182-1186.	
	GN	PANDEYA, S.N. et al., Synthesis of Some New Amidine Derivatives As Potent Hypoglycemic Agents, Pharmacological Research Communications, Vol. 17, No.8 (1985) pp. 699-709.	
	GO	PAULY, R.P. et al., Inhibition of Dipeptidyl Peptidase IV (DPIV) in Rat Results in Improved Glucose Tolerance, Regulatory Peptides Vol. 64, Issues 1-3 (1996) p. 148.	
	GP	PEDERSON, RAYMOND A. et al., Improved Glucose Tolerance in Zucker Fatty Rats by Oral Administration of the Dipeptidyl Peptidase IV Inhibitor Isoleucine Thiazolidine, Diabetes, Vol. 47 (1998) pp.1253-1258.	
	GQ	PILLAI, SREEKUMAR et al., Effects of ATP, Vanadate, and Molybdate on Cathepsin D-catalyzed Proteolysis, The Journal of Biological Chemistry, Vol. 280, No. 14 (1985) pp. 8384-9.	
	GR	PODANYI, BENJAMIN et al., Nitrogen Bridgehead Compounds. 62. Conformational Analysis of 6, 7, 8, 9-Tetrahydro-4H-pyrido[1,2-a]pyrimidin-4-ones and Their Methyl Derivatives by NMR Spectroscopy, J. Org. Chem. 51 (1985) 394-399.	
	GS	POJE, M. et al., Diabetogenic action of alloxan-like derivatives of uric acid, Experientia 36 (1980) pp. 78-9.	
	GT	POJE, M. et al., Oxidation of Uric Acid. 4. Synthesis, Structure, and Diabetogenic Action of 5-Imino-2,4,6-(1H,3H,5H)-pyrimidinetrione Salts and Their Alloxan-like Covalent Adducts, J. Med. Chem. 26 (1983) 861-4.	
	GU	POLACEK, I. et al., Hypoglycemic Activity of Amine Derivatives, Arzneim.-Forsch./ Drug Res. 28 (1978), 791-93.	
	GV	PRIDAL, L. et al., Glucagon-Like Peptide-1(7-37) Has a Larger Volume of Distribution Than Glucagon-Like Peptide1(7-36)amide in Dogs and is Degraded More Quickly in Vitro by Dog Plasma, European Journal of Drug Metabolism and Pharmacokinetics, Vol. 21 (1995), pp. 51-59.	

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	GW	RAM, VISHNU JI et al., Synthesis and Antihyperglycemic Activity of Suitably Functionalized 3H-quinazolin-4-ones, Bioorganic & Medicinal Chemistry 11 (2003), pp. 2439-2444.
	GX	SAWYER, JAMES H. et al., Pyrido[1,2-a]pyrimidinium Salts. Part 1. Synthesis from 2- Aminopyridines and Interconversion with 2-(2-Acetylvinylamino) pyridines, J.C.S. Perkin I (1972), 1138-1143.
	GY	SAXENA, A.M. et al., Mode of action of three structurally different hypoglycemic agents: A comparative study, Indian Journal of Experimental Biology, Vol. 34 (1996), pp. 351-355.
	GZ	SEDO, ALEKSI et al., Dipeptidyl peptidase IV-like molecules: homologous proteins or homologous activities? Biochimica et Biophysica Acta 1550 (2001), pp. 107-116.
	HA	SEKIYA, T. et al., Pyrimidine derivatives. III (1) Synthesis of hypoglycemic 4-alkoxy-2-piperazino-activity of 6-polymethylenepyrimidines, Eur. J. Med. Chem. (1982), 75-79.
	HB	SENTEN, KRISTEL et al., Development of Potent and Selective Dipeptidyl Peptidase II Inhibitors, Bioorganic & Medicinal Chemistry Letters 12 (2002) pp. 2825-2828.
	HC	SETH, M. et al., Syntheses of 2-Substituted & 2,3-Distributed 4(3H)-Quinazolones, Indian Journal of Chemistry, Vol. 14B (1975), 536-540.
	HD	SHIMAZAWA, RUMIKO et al., Novel Small Molecule Nonpeptide Aminopeptidase N Inhibitors with A Cyclic Imide Skeleton, J. Enzyme Inhibition, Vol. 14 (1999) pp. 259-275.
	HE	SHISHEVA, ASSIA, et al., Insulinlike Effects of Zinc Ion in Vitro and in Vivo; Preferential Effects on Desensitized Adipocytes and Induction of Normoglycemia in Streptozocin-Induced Rats, Diabetes, Vol. 41 (1992), pp. 982-988.
	HF	SINYAK, R. S. et al., Synthesis and Biological Properties of Derivatives of 4-Hetarylmercaptoquinazoline, Translated from Khimiko-farmatsevticheskii Zhurnal, Vol. 20, No. 2, pp 168-171 (1986), pp. 103-105.
	HG	SOKAL, JOSEPH E., Basal Plasma Glucagon Levels of Man, Journal of Clinical Investigation, Vol. 46, No.5 (1967) pp. 778-785.
	HH	SRIVASTAVA, P.P. et al., Efficacy of Centipiperalone in Combination With Biguanide & Sulfonylurea, Indian Journal of Experimental Biology, Vol. 21 (1983), pp. 390-392.
	HI	TANAKA, KEIJI et al., Vanadate Inhibits the ATP-Dependant Degradation of Proteins in Reticulocytes Without Affecting Ubiquitin Conjugation, The Journal of Biological Chemistry, Vol. 259, No. 4 (1983), 2803-2809.
	HJ	VILLHAUER, EDWIN B. et al., DPP-IV Inhibition and Therapeutic Potential, Annual Reports in Chemistry 36 (2001), 191-200.
	HK	VILLHAUER, EDWIN B. et al., 1-[[(3-Hydroxy-1-adamantyl)amino]acetyl]-2-cyano-(S)-pyrrolidine: A Potent, Selective, and Orally Bioavailable Dipeptidyl Peptidase IV Inhibitor with Antihyperglycemic Properties, J. Med. Chem. 46 (2003), pp. 2774-2789.

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HL	WELLS, CAROL L. et al., Role of Anaerobic Flora in the Translocation of Aerobic and Facultatively Anaerobic Intestinal Bacteria, Infection and Immunity, Vol. 55, No. 11 (1987) pp. 2689-94.	
HM	WIEDEMAN, PAUL E. et al., Dipeptidyl peptidase IV inhibitors for the treatment of impaired glucose tolerance and type 2 diabetes, Current Opinion in Investigational Drugs, Vol. 4, No. 4 (2003), pp. 412-420.	
HN	YASUDA, NOBUYUKI et al. Enhanced Secretion of Glucagon-Like Peptide 1 by Biguanide Compounds, Biochemical and Biophysical Research Communications 298 (2002), pp. 779-784.	
HO	YUEN, V.G. et al., Acute and Chronic Oral Administration of Bis(maltolato)oxovanadium(IV) in Zucker Diabetic Fatty (ZDF) Rats, Diabetes Research and Clinical Practice 43 (1999), pp. 9-19.	
HP	ZANDER, METTE, et al., Additive Glucose-Lowering Effects of Glucagon-Like Peptide-1 and Metformin in Type 2 Diabetes, Diabetes Care, Vol. 24, No. 4 (2001) pp. 720-725.	
HQ	ZHANG, ANQI et al., Vanadate Stimulation of Insulin Release in Normal Mouse Islets, The Journal of Biological Chemistry, Vol. 266, No. 32 (1991), pp. 21649-56.	

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